

**Amendments to the Claims:**

Please amend the claims as follows:

1. (currently amended) A method for sequence-specific identification, separation and quantitation of polynucleotide fragments in ~~amplification~~ of a population of polynucleotides comprising:
  - (a) reverse transcribing an RNA population to provide a double-stranded cDNA population;
  - (b) digesting said cDNA population with one or more restriction endonucleases having a degenerate recognition or cleavage sequence, wherein said restriction endonuclease is a three- to eight-base cutter and wherein the degenerate recognition or cleavage sequence is represented by the formula of  $N^m$ , where N is the extent of degeneracy, and m is the number of degenerate bases, and wherein for at least one of said restriction endonucleases N is 2-4 and m is 1-5, to produce restriction fragments having  $N^m$  different single-stranded overhangs for each restriction endonuclease;
  - (c) ligating said restriction fragments to a series of adapters lacking restriction endonuclease sites, each adapter having a sequence complementary to one of said overhangs such that restriction fragments having identical overhangs are ligated to the same adapter; [[and]]
  - (d) amplifying said restriction fragments for no more than 25 cycles with a primer comprising a detectable label; and
  - (e) detecting and quantifying said polynucleotide fragments.
2. (original) The method of claim 1 wherein for at least one of said restriction endonucleases m is 2, 3 or 4.
3. (original) The method of claim 1 wherein said restriction endonuclease comprises a four-base cutter.
4. (original) The method of claim 1 further comprising digesting the restriction fragments obtained in (b) with one or more further restriction endonucleases producing restriction fragments with single-stranded overhangs different from those produced in (b).
5. (original) The method of claim 4 further comprising ligating the single-stranded overhangs produced by the digesting of claim 4 to a series of adapters each adaptor having a sequences complementary to one of said overhangs.
6. (original) The method of claim 1 wherein said restriction fragments of (d) are amplified by the polymerase chain reaction (PCR) to produce PCR products.
7. (original) The method of claim 6 wherein said adapters provide priming sites for said polymerase chain reaction.
8. (original) The method of claim 6 further comprising detecting the PCR products.
- 9-23. (cancelled)